

Remarks

Claims 14-36 are pending in the subject application and are currently before the Examiner. Favorable consideration of the pending claims is respectfully requested.

Claims 14-36 are rejected under 35 U.S.C. § 112, first paragraph, as nonenabled by the subject specification. The Office Action argues that the as-filed specification fails to enable the claimed invention and that undue experimentation would be required to practice the invention as claimed. Applicant respectfully asserts that the claims are enabled by the subject specification and traverse the rejection.

When rejecting a claim under the enablement requirement of section 112, the examiner bears the “initial burden of setting forth a reasonable explanation as to why [he/she] believes that the scope of protection provided by [the] claim is not adequately enabled by the description of the invention provided in the specification.” *In re Wright*, 999 F.2d 1557, 1562, 27 USPQ2d 1510, 1513 (Fed. Cir. 1993). Further, to object to a specification on the grounds that the disclosure is not enabling with respect to the scope of a claim sought to be patented, the examiner must provide evidence or technical reasoning substantiating those doubts (*Id.*; and M.P.E.P. Section 2164.04) and without a reason to doubt the truth of the statements made in the patent application, the application must be considered enabling. *In re Wright*, 999 F.2d 1557, 1562, 27 USPQ2d 1510, 1513 (Fed. Cir. 1993); *In re Marzocchi*, 439 F.2d 220, 223, 169 U.S.P.Q. 367, 369 (C.C.P.A. 1971).

Applicant respectfully submits that the as-filed application teaches the specific types of interferons that are to be administered to an individual with SARS, the amounts of interferons and ribavirin that are to be administered to individuals with SARS and the modes and/or timing for the administration of the interferons and ribavirin (see as-filed specification at pages 7-8 and 13-15). Thus, adequate teaching exists within the as-filed specification as to how one skilled in the art is to practice (use) the claimed invention. Thus, contrary to the arguments proffered in the Office Action at page 3, adequate teaching as to the identity of the interferon to be administered as well as the routes of administration and dosages are provided in the as-filed specification. Applicant further notes that the Office Action also argues that post-filing date literature suggests that neither IFN nor IFN/anti-viral treatments are effective for SARS infections and that this supports a finding that “the type and amount of experimentation regarding developing effective IFN therapies now verges on

non-routine; similarly, co-administration of an additional, effective antiviral with IFN awaits development of effective anti-SARS-CoV therapeutics.” However, no supporting evidence for these assertions is provided in the Office Action. As such, Applicant respectfully submits that the Office Action fails to provide any evidence supporting the arguments that the instant application is not enabled and that the Examiner has not met his burden of establishing that the claimed invention is not enabled. Accordingly, reconsideration and withdrawal of the rejection is respectfully requested.

Claims 14, 17, 18, 25-29, 32, and 33 are rejected under 35 U.S.C. § 103(a) as obvious over Higgins *et al.* in view of Ksiazek *et al.* (2003), Arnason (1999), and Weinstock-Guttman *et al.* (2000). The Office Action states that Higgins *et al.* teach use of purified, recombinant human interferon to prevent or treat the human coronavirus, 229E. Ksiazek *et al.* teach that a coronavirus, SARS-CoV, is the etiologic agent of SARS. Arnason teach that IFN- β is effective if administered subcutaneously while Weinstock-Guttman *et al.* teach that it is also effective if administered intramuscularly. Applicant notes that the teachings of Weinstock-Guttman *et al.* and Arnason *et al.* are directed to the use of interferon- β for the treatment of multiple sclerosis and its subcutaneous or intramuscular administration. Claims 15, 16, 19, 20, 30, 31, and 34-36 are rejected under 35 U.S.C. § 103(a) as obvious over Higgins *et al.* (1983), Ksiazek *et al.* (2003), Arnason (1999), and Weinstock-Guttman *et al.* (2000) as applied to claims 14, 17, 18, 25-29, 32, and 33 above in further view of Albrecht (WO 99/151943, 1999). The Office Action states that an ordinarily skilled artisan would have been motivated to use interferon in combination with ribavirin for the treatment of SARS because no treatment for SARS was available at the time and interferon/ribavirin had previously been used to treat another RNA virus. Applicant respectfully asserts that the claimed invention is not obvious over the cited references. Accordingly, reconsideration and withdrawal of the rejection under 35 U.S.C. § 103(a) is respectfully requested.

Claims 21-24 are rejected under 35 U.S.C. § 103(a) as obvious over Higgins *et al.* (1983), Ksiazek *et al.* (2003), Arnason (1999), Weinstock-Guttman *et al.* (2000), and Albrecht (1999) as applied to claims 14-20 and 25-36 above, in further view of Chang *et al.* (U.S. Patent No. 5,723,125), Higgins *et al.*, Ksiazek *et al.*, Arnason, Weinstock-Guttman *et al.*, and Albrecht. The Office Action indicates that an ordinarily skilled artisan would have been motivated to use interferon/immunoglobulin chimeric protein for the treatment of SARS because no treatment for

SARS was available at the time and interferon/IgG had previously been used to treat another RNA virus. Applicant respectfully asserts that the claimed invention is not obvious over the cited references and traverses each of the rejections on the following grounds.

As the Patent Office is aware, one skilled in the art must have a reasonable expectation of success in carrying out the claimed method. *See Medichem*, 437 F.3d at 1164; *Noelle v. Lederman*, 355 F.3d 1343, 1351-52 (Fed.Cir.2004); *Brown & Williamson Tobacco Corp. v. Philip Morris, Inc.*, 229 F.3d 1120, 1121 (Fed.Cir.2000). In the case of the rejections of record, it is respectfully submitted that the teachings relied upon in establishing the rejection would not have provided one of ordinary skill in the art with a reasonable expectation of successfully carrying out the claimed invention, particularly in view of the arguments proffered in the enablement rejection issued in this matter. However, it is also noted that a post-filing date reference provides evidence that the claimed methods of treatment provide unexpected results. As noted in *De Albuquerque et al. (J. Virol.*, 2006, 80:10384-10394), a copy of which is attached hereto, treatment of animals in a model of SARS resulted in improved survival of animals (see Abstract). Accordingly, it is respectfully submitted that the claimed invention is not obvious over the cited combination of references and reconsideration and withdrawal of the obviousness rejections of record is respectfully requested.

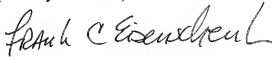
Applicant expressly reserves the right to pursue the invention(s) disclosed in the subject application, including any subject matter canceled or not pursued during prosecution of the subject application, in a related application.

Applicant believes that the currently pending claims are in condition for allowance, and such action is respectfully requested.

The Commissioner is hereby authorized to charge any fees under 37 CFR § 1.16 or 1.17 as required by this paper to Deposit Account No. 19-0065.

Applicant invites the Examiner to call the undersigned if clarification is needed on any of this response, or if the Examiner believes a telephonic interview would expedite the prosecution of the subject application to completion.

Respectfully submitted,



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Attachment: De Albuquerque *et al.*, 2006

Murine Hepatitis Virus Strain 1 Produces a Clinically Relevant Model of Severe Acute Respiratory Syndrome in A/J Mice

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Severe acute respiratory syndrome (SARS) is a life-threatening infectious disease which has been difficult to study and treat because of the lack of a readily available animal model. Intranasal infection of A/J mice with the coronavirus murine hepatitis virus strain 1 (MHV-1) produced pulmonary pathological features of SARS. All MHV-1-infected A/J mice developed progressive interstitial pneumonitis, including dense macrophage infiltrates, giant cells, and hyaline membranes, resulting in death of all animals. In contrast, other mouse strains developed only mild transitory disease. Infected A/J mice had significantly higher cytokine levels, particularly macrophage chemoattractant protein 1 (MCP-1/CCL-2), gamma interferon, and tumor necrosis factor alpha. Furthermore, FGL2/fibronectin mRNA transcripts and protein and fibrin deposits were markedly increased in the lungs of infected A/J mice. These animals developed a less robust type I interferon response to MHV-1 infection than resistant C57BL/6J mice, and treatment with recombinant beta interferon improved survival. This study describes a potentially useful small animal model of human SARS, defines its pathogenesis, and suggests treatment strategies.

Severe acute respiratory syndrome (SARS) was first diagnosed in China in November 2002 and quickly spread to Hong Kong, Singapore, Vietnam, Taiwan, and Canada (15, 20, 42). SARS was documented in approximately 8,000 persons globally and resulted in over 700 deaths. Outbreaks in research laboratory personnel and the recent discovery of a new reservoir of the etiologic virus argue that further outbreaks will arise (21).

The causative agent, SARS coronavirus (SARS-CoV), best fits within group 2 coronaviruses, which include the mouse hepatitis viruses (MHV) (6, 12, 15). The clinical severity of SARS varied considerably, presumably because of the genetic diversity of the host immune response (26). Considerable variation among SARS patients with respect to circulating viral load and patterns of SARS-CoV-evoked cytokine responses have been reported, although serum levels of gamma interferon (IFN- γ), IL-10, CXCL10 (IP-10), CCL5 (RANTES), and CXCL8 (IL-8) were elevated in most patients (54). However, the underlying pathogenetic mechanisms have not been clearly elucidated.

At present, no effective therapeutic strategies have been developed for SARS. Treatments initially used during the last outbreak of SARS included ribavirin and corticosteroids (38, 46). Subsequently, hyper-immune globulin, protease inhibitors, and IFNs were considered as alternative options for treating

SARS patients (57). Two groups, including our own, showed potential effectiveness for IFN in SARS (2, 27). In one study, IFN- α was shown to inhibit viral replication in vitro and to ameliorate disease in vivo. In a second study, our group demonstrated the potential efficacy of IFN- α in the treatment of SARS-infected patients (27).

Coronaviruses are known to cause a broad spectrum of diseases in animals, including pneumonia, hepatitis, nephritis, enteritis, and encephalitis. The type and severity of disease are influenced by the age and genetic background of the host, the route of infection, and the coronavirus strain (58). Members of the *Coronaviridae* family share replicative and transcriptional features. In humans, coronaviruses were thought to only cause upper respiratory infections and have only been rarely implicated in lower respiratory tract infections (33). Unlike influenza virus, human respiratory coronaviruses, other than SARS-CoV, do not cause epithelial cell necrosis. Thus, the SARS-CoV must have distinct characteristics leading to its unique pathogenicity.

MHV has served as a model for dissecting the viral and immunologic determinants of coronavirus disease (3, 28, 41, 53, 55). Extensive analysis of A59/JHM chimeric viruses has demonstrated that, while the spike gene is a major determinant of tropism and virulence in the central nervous system (39, 40), other viral genes play major roles in determining the ability to infect the liver (34) as well as the type and extent of both innate and T-cell responses induced (44). Infection of mice with recombinant viruses in which the replicase genes of JHM and A59 have been exchanged demonstrates that the presence of 3' structural genes rather than the replicase determines the dif-

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ferences in tropism and virulence between these strains (23). We have demonstrated that the macrophage prothrombinase FGL2/fibrinogen is an important determinant of disease in MHV-3-induced fulminant hepatitis and that the coronavirus nucleocapsid gene mediates much of its effect by inducing FGL2/fibrinogen (30, 36, 37). The MHV E gene is an inducer of apoptosis (1), the M gene of TGEV coronavirus is an inducer of IFN- α (22), and expression of the HE protein enhances the neurovirulence of a virus expressing the MHV-JHM spike (18). Resistance to MHV has been shown to involve both the host innate and adaptive immune systems (29, 47, 53, 56). Animals susceptible to MHV-3 generate an early increased proinflammatory response and predominant Th2 cytokine profile leading to activation of coagulation and tissue necrosis, whereas resistant animals generate a predominant Th1 immune response leading to production of cytotoxic T-lymphocyte and protective B-cell responses (4, 35, 41). In a comparison of a highly neurovirulent recombinant JHM with the weakly neurovirulent A59, the high lethality of the former is associated with an inability to induce a robust CD8T cell response (44). These studies serve as an excellent background for the study of a SARS model in mice.

None of the reported models of SARS produce lung pathology similar to that seen in humans and thus are best suited to studying the inhibitory effects of antiviral agents. In this report, we demonstrate that MHV-1 infection of inbred strains of mice produces a lethal SARS-like disease in A/J mice with features similar to those found in human patients, whereas other mouse strains including C57BL/6J mice fully recover. We further define differences in cytokine and type I IFN responses in resistant and susceptible mice and confirm a role for IFN in treatment for the viral disease.

MATERIALS AND METHODS

Mice. Female BALB/c, C57BL/6J A/J, and C3H/He mice 6 to 8 weeks of age (Charles River Laboratories and Jackson Laboratories) were maintained in microisolated cages and housed in the animal colony at Toronto General Hospital, University of Toronto, and fed standard lab chow diet and water ad libitum. All protocols were approved by the Animal Welfare Committee.

Virus. Parental viruses MHV-3, MHV-A59, MHV-JHM, MHV-S, and MHV-1 were obtained from the American Type Culture Collection, Manassas, VA. Viruses were first plaque purified and then expanded in murine J772A cells. Supernatants were collected and subsequently stored at -80°C until used. Mice were infected with 5,000 PFU intranasally unless otherwise stated.

Cells. The origins and growth of 17C-1, L2, and L929 cells have been previously described (48).

Viral infection and IFN- β treatment. All viral infection studies were performed in a viral infection room. Mice were anesthetized by intraperitoneal injection with 0.2 ml (or 10% of their body weight) of pentobarbital diluted in normal saline. Immediately, mice received an intranasal inoculation of 5×10^5 PFU MHV-1 in 50 μl ice-cold Dulbecco's modified Eagle's medium. The virus (5×10^5 PFU) was instilled into the nares, and mice were observed until the virus was inhaled. Mice were monitored daily for symptoms of disease, including ruffled fur, tremors, and lack of activity. Additionally, three groups of 10 A/J mice in each were infected intranasally with MHV-1 and either mock treated with phosphate-buffered saline (PBS) or treated with IFN- β . 10 mice received 10^5 IU of IFN- β 24 h prior to virus infection, 10 mice received 10^5 IU of IFN- β 12 h post-virus infection, and 10 mice received PBS at 12 h postinfection. IFN- β and PBS were administered by intraperitoneal inoculation.

Tissue isolation. Mice were sacrificed on days 2, 8, 14, and 21 postinfection. Blood was collected via cardiac puncture, and serum was stored at -80°C . Liver, lung, heart, kidney, spleen, and brain were collected at the times indicated for histology, viral titers, and molecular analysis. Samples denoted for histology were fixed in 3 ml of 10% formalin. Histology samples were processed by standard

methods. Samples for viral titers were snap-frozen in prelabelled cryovials in liquid nitrogen and subsequently stored at -80°C .

Viral titers. Samples were homogenized in 10% ice-cold DMEM utilizing a Polytion homogenizer (Fisher Scientific, Whittby, Ontario, Canada). Viral titers were determined in L2 cells as described previously (24).

Western blot analysis. Lung homogenates from each mouse strain were prepared for Western immunoblot analysis. Samples were processed for Western immunoblots as described previously (58). Briefly, cell lysates were resolved by sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and resolved proteins were transferred to membranes and then immunoblotted with antibody to MHV-1 nucleocapsid protein and visualized using chemiluminescence. β -Actin served as a loading control.

RNA extraction and cDNA synthesis. Animal tissues snap-frozen at -80°C were thawed, and then RNA was extracted using Invitrogen TRIzol reagent, as per the manufacturer's instructions. cDNA was synthesized using 1 μg RNA in the presence of random primers and Moloney murine leukemia virus reverse transcriptase (Invitrogen, Burlington, Ontario, Canada) according to the manufacturer's protocol.

Real-time PCR for detection of type I IFNs and FGL2. Reaction components were obtained from the LightCycler FastStart DNA Master SYBR green $\text{PL}^{1.2}$ kit (Roche). The LightCycler instrument (Roche) and corresponding software were used for all reactions. The reaction conditions used are as follows. The PCR was performed in a final volume of 20 μl containing 1 \times Master SYBR green $\text{PL}^{1.2}$ 1 buffer, various amounts of primer, and 5 μl of template cDNA (concentration, 100 ng/ml). The following primers and reaction conditions were used for real-time PCR: IFN- α , 5' ATGGCTAGGCTCTGTCGTTTC 3' and 5' TCA TTCTCTCTCTCAGCTCTC 3'; IFN- α , 5' AAAGGGGAGGCTCTCTCAT 3' and 5' TGCTTCTCTCTGATGTCGA 3'; IFN- α , 5' CATGATCTAGT AATGATGAGCTACTACT 3' and 5' TCAAGAGGAGGCTCTCTGATCAG 3'; IFN- α , 5' CAAAGCTCTGTGTGATGCA 3' and 5' ACTCTCTGCTCAAT CTTGCA 3'; IFN- α , 5' CCTCTCTCTCTGATGCTCT 3' and 5' CAGT TCTCTCATCCG 3'; IFN- α , 5' CTGCAAGATTTGATGAGGAAGAG AAGCT 3' and 5' ATAAAGCAACAAATAAATAACAAATAGGTGGC 3'; IFN- α , 5' AGCAAAATGGAGAGAACTGTCTAGA 3' and 5' ATGACCA TGTCTAATTAAGTCTT 3'; IFN- α , 5' ACACAGACTTACCAACATGACA ACAGGTGGATCTCTACGC 3' and 5' GTTAGAATTTTCACTTTG AGATTCTGTGATTCCTCTG 3'; Fgl2, 5' GTCAACGCGGTTCAG ATCT 3' and 5' TGTAGTCCCACTGCTGCTC; and hypoxanthine phosphoribosyltransferase (hprt), 5' CAAGCTGTGCTGTGAAAGAGA 3' and 5' TGAAGTACTCATATAGTCAAGGGCATATC 3'. The annealing temperatures and primer concentrations for the various genes utilized in real-time PCR analysis are as follows: IFN- α , 68°C and 10 pmol; IFN- α , 65°C and 10 pmol; IFN- α , 65°C and 10 pmol; IFN- α , 65°C and 10 pmol; IFN- α , 45°C and 10 pmol; IFN- α , 45°C and 10 pmol; IFN- β , 60°C and 20 pmol; Fgl2, 60°C and 10 pmol; and hprt, 65°C and 20 pmol.

Cytokine analysis. Serum cytokine levels were assayed using cytometric bead array kits (BD Biosciences, Mississauga, Ontario, Canada): a Th1/Th2 cytokine array kit detects IL-2, IL-4, IL-5, tumor necrosis factor α (TNF- α), and IFN- γ and an inflammation kit which detects IL-6, IL-10, CCL2 (MCP-1), IFN- γ , TNF- α , and IL-12p70. Serum samples were processed as per the manufacturer's instructions and analyzed using a Becton Dickinson FACScan flow cytometer.

Immunohistochemistry and in situ hybridization. (i) Double-staining immunohistochemistry to detect fgl2 and fibrin. Immunohistochemical staining for detection of the fgl2 prothrombinase was used to assess fgl2 prothrombinase expression in lung tissue. Tissues were fixed with 100% methanol prior to immunostaining. A polyclonal antibody to fgl2 prothrombinase was produced in rabbits by repeated injections with a 14-amino-acid hydrophilic peptide (CKLQADHIDHDPGGR) from exon 1 of the fgl2 gene, which had been coupled to keyhole limpet hemocyanin. Antibody was purified by affinity columns, and thereafter, tissue or cultured cell slices were incubated with antibody (20 $\mu\text{g}/\text{ml}$ in PBS). For fibrin detection, a rabbit antifibrinogen antibody (DakoCytomation, Carpinteria, Calif.) was utilized. Dual staining for fgl2 and fibrin was performed using a Vectastain ABC kit (Vector Laboratories, Burlingame, Calif.) with the second antibodies labeled with alkaline phosphatase or horseradish peroxidase, respectively. Subsequently, sections were incubated with immunoperoxidase-conjugated goat immunoglobulin G (6.7 $\mu\text{g}/\text{ml}$). Sections were photographed with a Leitz Laborlux fluorescence S microscope.

(ii) Immunohistochemical staining for the infiltrating cells in lung tissue. Infiltrating cells in lung tissue from A/J mice were characterized both preinfection (day 0) and on days 2 and 8 post-MHV-1 infection by indirect immunohistochemistry using a standard avidin-biotin complex (ABC) immunoperoxidase method. A rat anti-mouse F4/80 antigen (Serotec, Raleigh, NC) at a

TABLE 1. Ability of strains of MHV to produce SARS lung pathology in BALB/cJ mice

Classical feature of SARS	Histopathology of strain ^a				
	MHV-3	MHV-A59	MHV-1	MHV-S	MHV-JHM
Congestion	—	+	+++	—	—
Edema	—	Few foci	—	—	—
Hyaline membrane	—	—	+	—	—
Interstitial thickening	Limited	+	+	+	+
Interstitial inflammation	+	+	Focal	+	+
Alveolar infiltrate	+	+	+	+	+
Bronchus	Normal	Mild peribronchial inflammation	Resolving peribronchial inflammation	—	—
Pattern of disease	Pneumonitis	Focal inflammation	SARS-like (reversible)	Pneumonitis	Normal

^a Slown is a summary of histopathological findings in BALB/cJ mice infected with different strains of MHV, including MHV-3, MHV-A59, MHV-1, MHV-S, and MHV-JHM. Scores: —, absent; +, mild; ++, moderate; +++, severe. Based on the findings, MHV-1 produces the histopathologic picture most compatible with human SARS.

1/100 dilution for detection of macrophages, a rabbit anti-human T-cell CD3 cross-reactive with mouse (Sigma, St. Louis, Mo.) at 1/400 dilution for T cells, a rat anti-mouse CD45R/B220 (BD Pharmingen, Mississauga, Ontario, Canada) at a dilution of 1/50 for B cells, and a rat anti-mouse neutrophil (Cedarlane, Mississauga, Ontario, Canada) at a dilution of 1/50 for neutrophils were used.

For morphometry, 10 high-power fields per slide were counted. Positive cells were stained positive with immunoperoxidase (brown).

In situ hybridization. The method employed for *in situ* hybridization has been described previously (5). A digoxigenin (DIG)-11-UTP (Roche)-labeled cDNA probe was prepared from a cloned 169-bp fragment of *nl2* cDNA, representing nucleotides 756 (ACTGTGACAA...) to 924 (... GAGTAAGGA). The DIG-UTP-labeled probe concentration was determined by immunometric reaction with chemiluminescent detection, and the probes were stored at -80°C. Tissue sections were deparaffinized in 100% xylene and 100% alcohol, followed by prehybridization in 50% formamide and 2× SSC (1× SSC is 0.15 M NaCl plus 0.015 M sodium citrate). The hybridization mixture consisted of 50% deionized formamide, 5% dextran sulfate, 250 µg of salmon sperm DNA per milliliter, and 2 µg of DIG-labeled cDNA probe per milliliter in 2× SSC. The hybridization mixture with the probe was denatured by heating in an 85°C water bath and added to tissue sections for hybridization at 42°C overnight. Hybridization was followed by application of 3% blocking reagent and incubation with polyclonal anti-DIG Fab conjugated to alkaline phosphatase. Sections were counterstained with methylene green and mounted with Permount (Fisher Scientific, Whitby, Ontario, Canada) for viewing.

Electron microscopy. (i) **Transmission electron microscopy sample preparation.** Electron microscopy was performed on lung samples stored at 4°C in fixative (2.5% glutaraldehyde, 3.2% paraformaldehyde, 1.0 M phosphate buffer, pH 7.2). Samples were postfixated with 1% osmium tetroxide diluted in phosphate buffer, followed by en bloc staining with 2.5% uranyl acetate. Samples were then dehydrated using a graded series of ethanol and treated with propylene oxide. Epoxy resin (Epon-Araldite) was used as the embedding material, and sections were cut on a Reichert Ultratrac E microtome and collected on 300 mesh copper grids. Grids were stained using uranyl acetate followed by Reynolds' lead citrate. Grids were analyzed with a Hitachi H7000 electron microscope at 75 kV.

(ii) **Immunogold electron microscopy.** Tissues were fixed in 4% paraformaldehyde and 0.1% glutaraldehyde in 0.1 M phosphate buffer, pH 7.4, and then mixed to 1-mm³ cubes in phosphate buffer, infused with 2.3 M sucrose prior to freezing in liquid nitrogen. Samples were then stored at -85°C for 48 h in a solution of absolute methanol containing 0.5% uranyl acetate. The samples were warmed to -20°C and then infiltrated with Lowry's HM20 (Marivic Services, Halifax, Nova Scotia, Canada) and embedded and polymerized in the cold with UV. Ultrathin sections were mounted on Formvar-coated nickel grids. The sections were incubated with antibody against MHV-N (nucleocapsid protein) followed by incubation with goat anti-murine immunoglobulin G-10-nm gold particle complexes (GF Healthcare, Piscataway, N.J.) and stained with uranyl acetate. The grids were examined using a JEOL JEM 1240 electron microscope (JEOL, Tokyo, Japan).

RESULTS

MHV-1 produces SARS-like pulmonary disease in BALB/cJ mice. In a first set of experiments, we examined the ability of a large number of murine coronavirus strains, including MHV-1, MHV-A59, MHV-JHM, MHV-S, and MHV-3 to produce SARS-like pathology (Table 1). BALB/cJ mice were initially studied as they are known to be highly susceptible to many MHV strains when infected by the intraperitoneal and intracerebral routes. BALB/cJ mice inoculated with 10⁵ PFU of MHV-1 intranasally developed pulmonary disease with partial characteristic features of human SARS, including interstitial pulmonary infiltrates, hyaline membranes, giant cells, congestion, and hemorrhage (Fig. 1a). In addition to diffuse pulmonary infiltrates, a few focal deposits of fibrin were seen around small arterial blood vessels and in alveolar spaces with entrapment of platelets. Changes were noted as early as 2 days postinfection and progressed to day 14. Clinically these mice became lethargic with rapid respiration, but all mice survived and pulmonary pathology resolved by day 21. Livers of MHV-1-infected BALB/cJ mice sacrificed on day 8 showed portal infiltrates which by day 14 progressed to portal-portal bridging (data not shown). In contrast, BALB/cJ mice inoculated intranasally with MHV-S and MHV-JHM developed neither liver nor lung pathology. Although MHV-3 and MHV-A59 produced pulmonary lesions, these were milder than those generated by MHV-1 and did not have the characteristics of lesions described in SARS-infected patients. In addition, MHV-A59- and MHV-3-infected mice all developed severe hepatic necrosis and died of liver failure by day 10, and thus these strains of MHV do not represent relevant models of SARS. Taken together, these studies suggested that MHV-1 produced a mild SARS-like disease in BALB/cJ mice, a mild but significant liver injury, but the disease was transitory and all of the infected mice survived.

MHV-1 infection of A/J mice produces clinical and pathological SARS-like disease with high mortality. Since the C57BL/6J, C3H/He, and A/J strains have previously been shown to exhibit various degrees of susceptibility to other coronaviruses, including MHV-3 and MHV-A59, we inoculated all of these strains of mice with MHV-1 intranasally. MHV-1-infected C57BL/6J mice developed only mild pulmonary disease

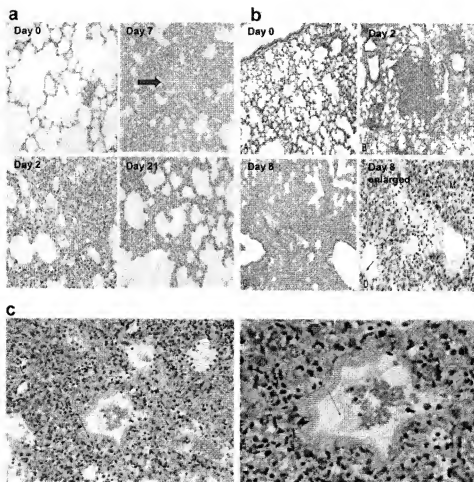


FIG. 1. Histopathology of MHV lung infection. (a) Histopathology of BALB/cJ mice infected with MHV-1 showing resolving interstitial pneumonitis. On day 2 postinfection, hemorrhage and inflammatory interstitial cell infiltrates are prominent. By day 7, hyaline membranes can be seen (arrow), and by day 21, pneumonitis is resolving (hematoxylin and eosin stain; magnification, $\times 200$). (b) Histopathology of MHV-1-infected A/J mice. The upper left panel represents day 0 postinfection, showing normal alveoli (hematoxylin and eosin stain; magnification, $\times 100$). The upper right panel represents 2 days post-MHV infection, showing mononuclear infiltrates with nodular formation (hematoxylin and eosin stain; magnification, $\times 100$). The lower left panel shows severe diffuse pneumonitis with consolidation on day 8 postinfection (magnification, $\times 100$). The lower right panel is a close-up of the lower left panel showing mononuclear cell infiltrates and hyaline membrane (arrow) (magnification, $\times 200$). (c) MHV-1-infected A/J mice on day 8 postinfection showing details of interstitial inflammation and hyaline membranes (arrow). The right panel is a higher magnification ($\times 500$) of the left panel ($\times 250$) showing further details of hyaline membranes.

similar to that seen in BALB/cJ mice by day 2 postinfection. The pulmonary disease progressed to day 8, but by day 14 postinfection, the pulmonary lesions improved and completely resolved (data not shown).

In contrast, A/J mice developed severe progressive pulmonary disease by day 2 post-MHV-1 infection and all MHV-1-infected A/J mice died within 7 to 10 days of infection. On day 2, patchy interstitial alveolar thickening and fluid accumulation in alveolar spaces (pulmonary edema) were prominent. At death, lungs showed severe interstitial pneumonitis with large areas of complete consolidation of the lungs. The interstitial inflammatory reaction included hyaline membranes, fibrin deposition, and heavy lymphocyte and macrophage infiltrates (Fig. 1b and c). By immunohistochemistry, infiltrating cells in MHV-1-infected lung tissue from A/J mice were predominantly macrophages and neutrophils on both day 2 and day 6 post-MHV-1 infection, as determined by morphometric anal-

ysis. T cells (CD3 positive) were also increased in the infiltrates, especially by day 6 postinfection. In contrast, B cells were relatively unchanged. (Fig. 2a and b).

Examination of the livers of MHV-1-infected A/J mice showed near normal histology to day 6, but on day 7 just prior to death, there was severe hepatic congestion compatible with lung and heart failure as described in humans (data not shown).

Following infection with MHV-1, C3H/St mice showed an intermediate pattern of resistance/susceptibility and surviving C3H mice went on to develop chronic pulmonary changes including fibrosis and bronchial hyperplasia (data not shown). Thus, A/J mice are highly susceptible to MHV-1-induced pulmonary disease when the virus is delivered intranasally, C57BL/6J are relatively resistant, and C3H mice have an intermediate susceptibility.

A/J, C3H/St, BALB/cJ, and C57BL/6J mice ($n = 10$ per

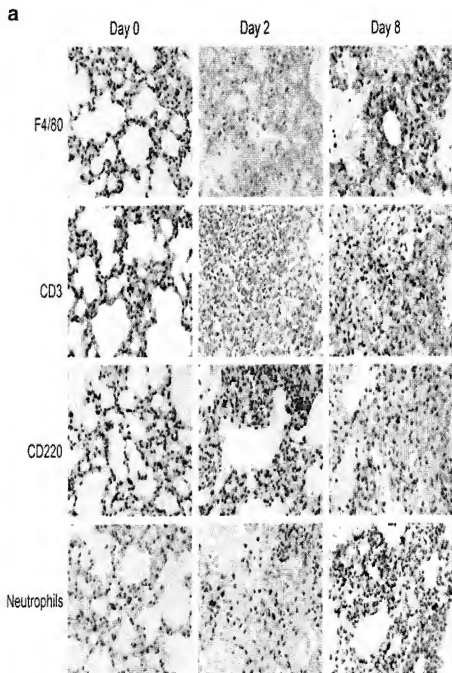


FIG. 2. (a) Immunoperoxidase stains of infiltrating cells in lung tissues from MHV-1-infected A/J mice. Columns represent days 0, 2, and 8. Rows represent different antibodies. Results are scored as follows: -, negative; +, mild positive; ++, moderate positive; +++, marked positive; +++++, most marked. (Top row) F4/80 macrophages. (Day 0) Lung showing normal tissue. Occasional resident macrophages in alveolar wall are lightly stained. No infiltrate was seen. Score, +/- (Day 2) Lung tissue showing moderate infiltrate of macrophages. Score, ++. (Day 8) Lung with consolidated heavy infiltrate of macrophages. Score, +++++. (Second row) CD3 T cells. (Day 0) Lung showing normal tissue with mild background staining. Score, -. (Day 2) Lung with scattering of positively stained T lymphocytes. Score, -. (Day 8) Lung consolidation and numerous positive T lymphocytes. Score, +++. (Third row) CD220 B cells. Lung shows occasional positive B lymphocytes. Score, +. (Bottom row) Neutrophils. (Day 0) Lung with no positively stained cells in this micrograph. Score, -. (Day 2) Lung infiltrate with a small number of neutrophils. Score, 1. (Day 8) Lung consolidation and numerous neutrophils. Score, +++. All micrographs are of immunoperoxidase-stained slides. Magnification, $\times 400$. (b) Morphometric analysis for specific cell lineage infiltration. Data are expressed on the y axis as the number (mean \pm standard deviation) of positive cells per high-power field (HPF). The number of days postinfection is given on the x axis. The lung infiltrate was a mixed-cell type with a marked increase in macrophages (F4/80), but increases in neutrophils and T cells (CD3⁺), especially by day 8 postinfection, were observed.

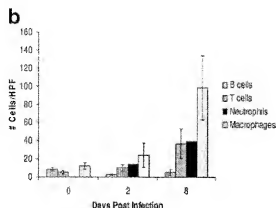


FIG. 2—Continued.

group) were inoculated intranasally with MHV-1 (5×10^3 PFU) and monitored closely for clinical signs of disease and survival (Fig. 3a). A/J mice, which are resistant to MHV-3 infection, developed clinical signs of disease within 48 h of intranasal inoculation of MHV-1 infection and succumbed within 8 days. In contrast, although BALB/cJ and C57BL/6J mice developed pulmonary disease, these animals cleared virus by day 14 and all mice survived. C3H mice developed an intermediate pattern of susceptibility with 40% mortality (Fig. 3a). A/J mice were then inoculated with increasing concentrations of MHV-1 (5×10^3 to 5×10^5 PFU) intranasally, and survival was monitored to day 21 (Fig. 3b). In dose-response studies, the 50% lethal dose (LD_{50}) for A/J mice infected with MHV-1 was calculated to equal 2.4×10^5 PFU using the method of Reed and Muench (43).

Expression of FGL2/fibrinogen is associated with pulmonary fibrin deposits. Fibrin deposition is a prominent feature

of SARS (7, 16). Our previous studies of coronavirus-induced fulminant hepatitis demonstrated that cell surface expression of the immune coagulant fgl2/fibrinogen led to organ-damaging microvascular thrombosis (25, 30). We therefore examined lung tissue from both susceptible A/J mice and resistant C57BL/6J mice infected with MHV-1 for the presence of FGL2 mRNA transcripts, protein, and fibrin (Fig. 4). As shown in Fig. 4a, fgl2 protein was expressed by inflammatory cells and type 1 pneumocytes in juxtaposition to deposits of fibrin. Furthermore, by in situ hybridization, FGL2 transcripts were detected in the cytoplasm of terminal alveoli and respiratory bronchioles (Fig. 4b). Also of interest, the bronchial epithelial cells are fused, forming giant cells, a hallmark of human SARS (Fig. 4b). Lungs from BALB/cJ and C57BL/6 mice had neither FGL2 nor fibrin deposits. Real-time reverse transcriptase PCR (RT-PCR) quantification of FGL2 transcripts demonstrated increased FGL2 mRNA expression in the lungs of A/J mice at 2 and 4 days postinfection, whereas FGL2 mRNA expression decreased in C57BL/6 mice post-MHV-1 infection (Fig. 4c).

Virus localization and viral burden in MHV-1-infected mice. A detailed examination of the lungs was made using transmission electron microscopy and immunogold electron microscopy on MHV-1 lung samples. On day 1 postinfection, virions were identified in the peripheral airways adjacent to capillaries in the alveolar walls (Fig. 5, left panel). A cluster of virions is seen at the arrow. The same particles are seen at higher magnification in the panel on the right. The virus particles occurred singly or in clusters. Individual particles were easily recognized by their uniform size and electron density. This result is important because it confirms that the virions reached the distal airways. There is also evidence of reaction to the viruses by the greatly enhanced vesicular activity noted in the capillary shown; it reflects the transport of fluid across the cell, which is greatly enhanced. It is a manifestation of edema formation,

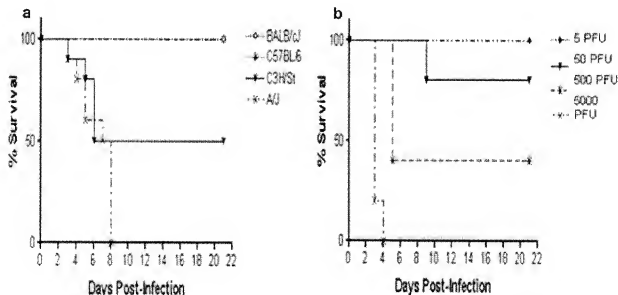


FIG. 3. Effect of MHV-1 on survival of inbred strains of mice. (a) Effect of MHV-1 infection on survival of BALB/cJ, C57BL/6J, A/J, and C3H/ST mice ($n = 10$ per group). Panel b shows the effect of increasing doses of MHV-1 on survival of A/J mice. Data were used to determine the LD_{50} by the method of Reed and Muench (43).

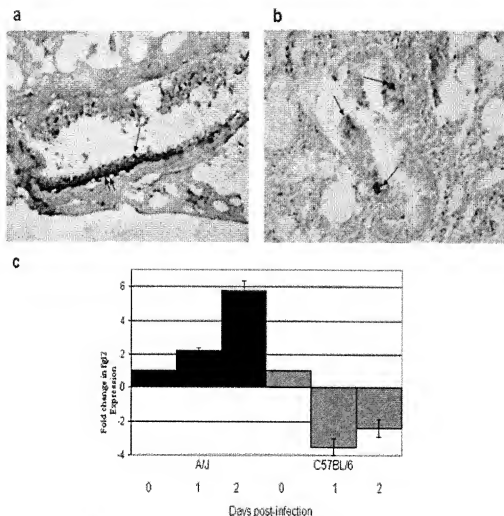


FIG. 4. Localization of fibrinogen-like protein (fgl2) and fibrin by immunohistochemistry and in situ hybridization in lung tissues from MHV-1-infected A/J mice 2 days postinfection. (a) By immunohistochemistry, bronchial epithelium is positive for fgl2 protein (blue staining, single arrow); dense fibrin deposits are seen along the wall of the bronchial (red staining, double arrow) (magnification, $\times 400$). (b) FGL2 mRNA transcripts are seen in bronchial lining cells (blue staining, single arrow); many are fused, forming giant cells (magnification, $\times 400$). (c) The increased number of FGL2 mRNA transcripts in lung of MHV-1-infected A/J mice was determined by real-time PCR. In contrast, no increase in FGL2 mRNA transcripts was seen in lung in C57BL/6J mice. Data represent the mean \pm standard deviation of three independent experiments done in triplicate.

which is an early reaction to the presence of the virus. Note also the neutrophil in the capillary lumen. By day 4, there was a notable infiltrate as seen also by light microscopy with lymphocytes, neutrophils, and macrophages. Electron microscopy showed extensive alveolar wall damage with fibrin, cell fragments, free red blood cells, virions, inflammatory cells, lymphocytes, macrophages, and neutrophils (not shown). By day 8, the lung was consolidated; the electron microscopic examination confirmed the predominance of macrophages, which often occurred in sheets and were the predominant cells in the infiltrate. Many cell fragments and red blood cells were also seen; alveolar walls were largely destroyed. A striking finding was that the macrophages were filled with viral particles. Measurement of the particles using the electron microscope point-to-point measurement software confirmed the correct measurement of the particles as expected

for coronavirus virions. To further confirm that they were indeed coronaviruses, immunoelectron microscopy was done, which confirmed their presence, their size, and their occurrence within macrophages. Note the presence of gold-labeled antibodies localized to the particles within macrophages. This is a very heavy infection of viruses; each macrophage is filled with virions. (Fig. 5)

Viral burden in lung, brain, spleen, and liver from both C57BL/6J and A/J mice was determined, using a standard plaque assay, as described previously (24). Virus was first detected in the lungs of A/J mice at 12 h postinfection, reached maximal levels by 48 to 72 h, and persisted at high levels until death of the animals on day 7 postinfection (Fig. 6a, left panel). Although virus was also recovered from the brain, liver, and spleen of MHV-1-infected A/J mice, it was detected later in the course of disease and at 4- to 5-log-fold-lower levels than

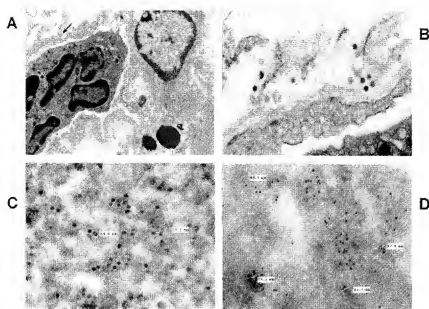


FIG. 5. Transmission electron microscopy (TEM [A to C]) and immunoelectron microscopy (IEM [D]) detection of virions in A/J lung tissues. (A) Virions (indicated by the arrow) are detectable in peripheral pulmonary airways; they are located outside the capillary wall of the lung on day 1. Viral particles are in close proximity to the plasma membrane of an endothelial cell within the alveolar wall. A neutrophil (darkly staining cell) is observed within the blood vessel. (B) Higher magnification of panel A. Virions are adjacent to endothelial cell basement membranes. In addition, vesicles are apparent in large numbers in endothelial cells of alveolar capillaries. By TEM (C), the small dense particles are virions are packed into lysosomes within a macrophage. Virion size (as indicated, ranging from 65 to 90 nm in diameter) corresponds to MHV. By IEM (D), 10-nm gold particles conjugated to anti-MHV nucleocapsid antibody are localized within pulmonary macrophages, confirming the viral nature of the particles. Virion diameters are 65 to 90 nm, typical of coronaviruses.

that seen in lung tissue (Fig. 6a, left panel). Virus was also detected in the lungs of disease-resistant C57BL/6J mice by 12 h postinfection, but at a lower titer than in A/J mice (Fig. 6a, right panel). Maximal levels of virus were reached by day 4, but no virus was detected by day 7 to 10 postinfection.

Lung tissues were examined for the presence of nucleocapsid protein by Western immunoblotting (Fig. 6b). In lung tissue from A/J mice, nucleocapsid protein was detected as early as 12 h postinfection and persisted to time of death of the animals on day 8 postinfection. Nucleocapsid protein was also detected in the lungs of C57BL/6J and BALB/cJ mice by day 2 postinfection and was detected even at 21 days postinfection at a time when infectious virus was undetectable by plaque assay.

Immune response to MHV-1: proinflammatory cytokine levels correlate with disease activity. In subsequent time course experiments, we examined the cytokine expression profiles in sera from MHV-1-infected A/J and C57BL/6J mice. Serum was collected from A/J and C57BL/6J mice both prior to MHV-1 infection and at various times postinfection and analyzed for IL-10, IL-6, IL-12p70, IFN- γ , TNF- α , and macrophage chemoattractant protein 1 (MCP-1/CCL2) (Fig. 6c). Notably, we observed marked elevation of IL-6, IL-10, IFN- γ , TNF- α , and CCL2 expression in infected A/J mice compared with infected C57BL/6J mice. In a separate set of experiments, sequential measurements of serum levels of TNF- α and IFN- γ were undertaken in both resistant and susceptible mice (Fig. 6d). At all time points postinfection, levels of both cytokines were higher in susceptible A/J mice. Levels of TNF- α peaked on day 6 postinfection, whereas levels of IFN- γ were highest at 2 days

postinfection. These results parallel observations in SARS patients.

IFN response and protective effects of IFN- β in MHV-1-infected A/J mice. To investigate the IFN response in the differentially susceptible strains, lung tissues from MHV-1-infected C57BL/6J and A/J mice were examined for gene expression for different IFN- α subtypes and IFN- β . Notably, levels of IFN induction, as measured by gene expression, were significantly lower in A/J than in the C57BL/6J mice (Fig. 7a). Transcriptional activation of the early response IFNs, IFN- β , and IFN- $\alpha 4$ was evident by 12 h postinfection, whereas gene expression for IFN- $\alpha 1$, IFN- $\alpha 2$, and IFN- $\alpha 5$ reached maximal levels at 24 h postinfection and then returned to basal levels by day 7 postinfection. No virus-inducible gene induction for IFN- $\alpha 9$, IFN- $\alpha 11$, or IFN- $\alpha 12$ was detectable in either mouse strain challenged with MHV-1.

Given our observation that A/J mice exhibited lower levels of type I IFN expression in the lung, we next examined the effect of administering exogenous IFN- β on A/J survival in response to MHV-1 infection. In this experiment, A/J mice were infected and either left untreated or treated with IFN- β . Two different IFN treatment regimens were employed: a single dose of 10^6 U/mouse at 24 h prior to virus infection or a single dose of 10^6 U/mouse 12 h post-virus challenge. As expected, the mice in the untreated group all succumbed to MHV-1 infection by day 5 postinfection with MHV-1 (Fig. 7b). Mice that received IFN- β 12 h post-virus inoculation exhibited an increased survival time, but all eventually succumbed to disease. Mice that were treated with IFN- β 24 h prior to viral

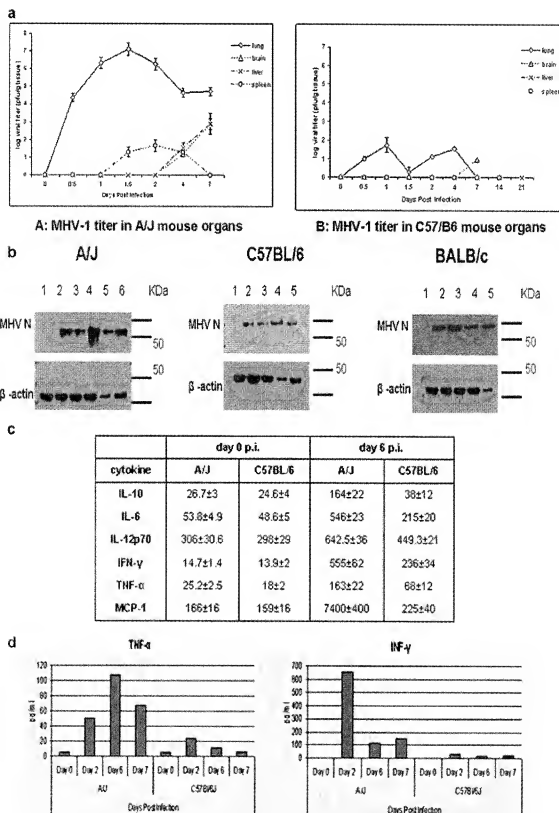


FIG. 6. Viral titers in tissues and cytokine responses. (a) Viral titers from lung (\circ), brain (Δ), liver (\times), and spleen (\square) from MHV-1-infected A/J mice and C57BL/6 mice at various times postinfection. (b) Western immunoblot detection of nucleocapsid protein in lungs. A/J mice: lane 1, day 0; lane 2, 12 h postinfection; lane 3, 24 h postinfection; lane 4, 4 days postinfection; lane 5, 6 days postinfection; and lane 6, 8 days postinfection.

challenge showed delayed onset of disease and an increased survival rate.

DISCUSSION

Although a number of models for SARS have been proposed, including SARS-CoV infection of mice (49), SARS-CoV infection of cats and ferrets (31), and SARS-CoV infection of nonhuman primates (13, 32), none of the models produce lung pathology or mortality similar to that seen in humans. Thus, these models can be utilized at best for evaluating the effectiveness of agents, including neutralizing antibodies, putative vaccines, or antiviral agents on viral replication. As reported herein, the MHV-1 mouse model produces a SARS-like clinical and pathological syndrome in which pathogenic mechanisms can be dissected.

MHV-1 was first isolated in Mill Hill, United Kingdom, and was reported to cause fatal hepatitis in the inbred P strain mice (10). Even in early studies, the pathogenicity of the virus was variable and critically dependent upon a number of immunomodulating variables, including coinfection with enterococci (11), Friend virus, or Moloney virus (9), and neurotropic and pneumotropic MHV-1 variants have been described (50, 52). Similarly, host age plays an important role in susceptibility to the virus; C57BL/6J, C3H, P, and Swiss Webster strains of mice have been noted to be very susceptible to intraperitoneal MHV-1 in the first 2 weeks of life, but susceptibility decreases as they age (10, 51, 52).

In the present study, we found that intranasal infection with MHV-1 produces a strain-dependent disease. The A/J mice, known to be resistant to the lethality of MHV-A59 and MHV-3 infection, all died following intranasal infection with MHV-1. Although BALB/cJ and C57BL/6J mice developed pulmonary disease, these animals cleared the virus by day 14 and survived. C3H mice developed an intermediate pattern of susceptibility with 40% mortality. The cell infiltrates seen in MHV-1-infected A/J mice were predominantly macrophages, with modest numbers of CD3⁺ T cells and neutrophils similar to that seen in human SARS (16). Peak viral titers were 10⁴-fold higher in susceptible A/J mice in comparison to resistant C57BL/6J mice, an observation confirmed by lung tissue electron microscopy in which more virions were present in A/J than C57BL/6J and BALB/cJ lung tissue cells. Interestingly, virions were seen largely in pulmonary macrophages rather than epithelial cells, consistent with our previous observations in MHV-3-induced fulminant hepatitis. These findings argue that host factors are critically important for the development of SARS-like disease and suggest that, as for other coronavirus diseases, innate immune cells are an important reservoir for infection. The finding that pulmonary gp2/fibronectin is strongly expressed in susceptible animals, but not in resistant ones, as

discussed below, is also consistent with a role for the innate immune system in acute disease.

Serum cytokines and chemokines were markedly elevated in susceptible A/J mice in comparison to those in resistant animals following MHV-1 infection. These findings are consistent with other respiratory diseases. For example, the role of proinflammatory cytokines in respiratory infections and acute respiratory distress syndrome has been previously documented (19, 45). Elevated levels of chemokines including CCL2 and CCL3 are seen in respiratory syncytial virus infections. Equally, elevations in IL-1 β , TNF- α , IFN- γ , and IL-12 have been reported in mice infected with human adenovirus (17). Patients with acute respiratory distress syndrome (ARDS) also have marked elevations in TNF- α and IL-6 (45). These elevated cytokines and chemokines may contribute to the immunopathology of SARS. Recently we and others have reported that serum concentrations of IFN- γ , IL-10, CXCL10, CCL5, and CXCL8 were elevated in SARS patients, especially in those patients with a poor outcome. In SARS patients, expression levels of IL-10, IFN- γ , and CXCL10 peaked within 4 days of peak viral titers, whereas IL-12p70, IL-4, and TNF- α were highest 5 to 7 days after peak viral loads. Similarly, during MHV-1 infection of A/J mice, high levels of IFN- γ and TNF- α were seen after peak viral titers were observed. The fact that corticosteroids ameliorated disease in some SARS patients is consistent with the hypothesis that an immunopathological mechanism may underlie the lung damage associated with SARS (54) as it is known that corticosteroids inhibit formation of proinflammatory cytokines. Furthermore, it has been reported that lower levels of both IL-6 and TNF- α are associated with better responses to steroid treatment in ARDS (14, 54).

We previously reported that MHV-3 induces FGL2, an inflammatory immune coagulant, which results in fibrin deposition and hepatic necrosis (24, 30). In the MHV-1 model of SARS, both FGL2 mRNA transcripts and fg12 protein were also seen in close association with deposits of fibrin in diseased lungs from A/J mice, suggesting that this inflammatory mediator may be contributing to the pathogenesis of SARS as well. Previously we reported that the nucleocapsid protein from strains of virus that caused massive hepatic necrosis, including MHV-3 and MHV-A59, induced transcription of FGL2. At present, we have no information whether the N gene from MHV-1 similarly induces FGL2 transcription. More recently, we have reported that IFN- γ can induce macrophage production of FGL2 and that, although TNF- α alone cannot induce FGL2 transcription, it synergizes with IFN- γ , leading to a marked enhancement in FGL2 transcription and subsequent thrombosis (8, 24, 30). In view of the findings of marked elevations of IFN- γ and TNF- α , both in the MHV-1-induced

BALB/cJ mice: lane 1, uninfected control; lane 2, 2 days postinfection; lane 3, 8 days postinfection; lane 4, 14 days postinfection; lane 5, 21 days postinfection. C57BL/6J mice: lane 1, uninfected control; lane 2, 2 days postinfection; lane 3, 8 days postinfection; lane 4, 14 days postinfection; lane 5, 21 days postinfection. Nucleocapsid protein was detected as a 50-kDa band. Measurement of β -actin levels was performed to ensure equal loading of protein. (c) Determination of inflammatory cytokines from uninfected and MHV-1-infected A/J and C57BL/6J lungs on day 0 or 6 postinfection. (d) Sequential measurements of serum levels of TNF- α and IFN- γ were undertaken in both resistant and susceptible mice. At all time points postinfection, levels of both cytokines were higher in susceptible A/J mice. Levels of TNF- α peaked on day 6 postinfection, whereas levels of IFN- γ were highest at 2 days postinfection.

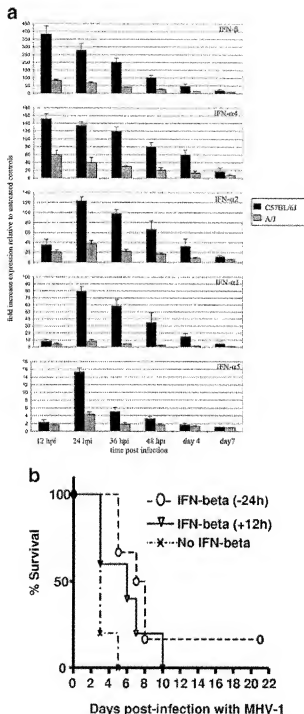


FIG. 7. Levels of type I interferon correlate with resistance to MHV-1. (a) Mice were infected intranasally with 10^5 PFU of MHV-1, and at each of the time points indicated, five mice were euthanized, their lung tissue was harvested, the RNA was extracted, and the cDNA was synthesized, as described in Materials and Methods. For each sample, reference (HPRT) and target (IFN- α 1, IFN- α 2, IFN- α 4, IFN- α 5, IFN- α 9, IFN- α 11, IFN- α 12, and IFN- β) quantitative RT-PCRs were performed. The data are presented as the change (fold) in expression relative to an untreated control sample \pm standard error. (b) A/J mice ($n = 10$ per group) were (i) treated with 10^5 IU of IFN- β 24 h prior to intranasal infection with 10^5 PFU of MHV-1 (○) or 12 h postinfection (inverted triangle) or (ii) treated with PBS at 12 h postinfection (×). Mice were monitored for survival once daily.

experimental model and in patients with SARS-CoV infection, this could have relevance to disease pathogenesis.

Based on the observation that intranasal infection with MHV-1 results in a dramatically more aggressive disease in A/J mice versus C57BL/6J mice, we examined whether the type and extent of the lung IFN response to infection might shape the outcome. A/J mice displayed significantly lower levels of type I IFN gene induction compared with C57BL/6J mice. Gene induction for IFN- α 9, IFN- α 11, and IFN- α 12 was absent in response to MHV-1 in both mouse strains. The kinetics of gene induction, namely early expression of IFN- β and IFN- α 4 and later gene expression of IFN- α 1, IFN- α 2, and IFN- α 5, agree with the sequential phosphorylation and activation of IRF3 followed induction and activation of IRF7. We next examined the effect of IFN- β treatment on the course of viral disease in A/J mice. The data revealed protective effects of IFN- β treatment. Mice that received IFN- β 24 h prior to virus infection exhibited a delayed onset of disease and increased survival. At the end of the study, mortality had decreased from 100 to 80% and 50% survival was extended from 3 to 8 days. Mice that received IFN- β 12 h post-virus infection also exhibited a delayed onset of disease, but survival was unaffected. The potential benefit of IFN treatment for respiratory coronavirus infections was reported in patients treated with IFN alfacon-1 during the recent SARS outbreak in Toronto. IFN alfacon-1-treated SARS patients displayed a more rapid recovery than patients treated with corticosteroids, in terms of time taken to resolve lung abnormalities, their oxygen saturation, their requirement for supplemental oxygen, and resolution of clinical measurements indicative of disease severity: lactate dehydrogenase and creatine kinase levels (29).

Although it is unclear whether SARS will continue to be a major human health problem, the isolation of SARS-CoV from Himalayan palm civets and bats suggests that an animal reservoir for SARS-CoV continues to exist (21). Furthermore, bat feces is used in traditional Chinese herbal medicines for treatment of asthma and kidney ailments and general malaise and additionally bat meat is a delicacy. Therefore, it is quite possible that additional SARS outbreaks will occur.

Thus, this model offers the potential to provide further insights into the pathogenesis of coronavirus-induced lung injury and the contribution of both the virus and host immune response. Furthermore, the data that we have generated in our animal model has great relevance to the pathogenesis of human SARS and support a role for IFN treatment as an effective therapy.

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